

(19)



Eur päisches Pat ntamt
European Patent Office
Office européen des brevets



(11) Publication number:

0 478 326 B1

(12)

EUROPEAN PATENT SPECIFICATION

(45) Date of publication of patent specification: **18.01.95** (51) Int. Cl.⁸: **B01J 13/02**

(21) Application number: **91308789.6**

(22) Date of filing: **26.09.91**

(54) Encapsulating method and products containing encapsulated material.

(30) Priority: **27.09.90 GB 9021061**

(43) Date of publication of application:
01.04.92 Bulletin 92/14

(45) Publication of the grant of the patent:
18.01.95 Bulletin 95/03

(84) Designated Contracting States:
BE CH DE FR GB LI NL

(56) References cited:
EP-A- 0 294 206
EP-A- 0 309 054

DATABASE/WPI, accession no. 79-04743B
[03], Derwent Publications Ltd, London,GB; &
JP-A-53 138 736 (RICOH K.K.)

DATABASE/WPI, accession no. 75-82558W
[50], Derwent Publications Ltd, London,

GB; & JP-A-50 034 516 (FUJI PHOTO FILM
K.K.)

(73) Proprietor: **QUEST INTERNATIONAL B.V.**
Hulzerstraatweg 28
NL-1411 GP Naarden (NL)

(72) Inventor: **Behan, John Martin**
Shermal,
Ball Lane,
Kennington
Ashford,
Kent (GB)
Inventor: **Ness, Jeremy Nicholas**
22 River Court,
Chartham
Canterbury,
Kent CT4 4JN (GB)
Inventor: **Perring, Keith Douglas**
14 Malvern Road
Ashford,
Kent TN24 7HS (GB)

(74) Representative: **Gura, Henry Alan et al**
MEWBURN ELLIS
York House
23 Kingsway
London WC2B 6HP (GB)

Note: Within nine months from the publication of the mention of the grant of the European patent, any person may give notice to the European Patent Office of opposition to the European patent grant. Notice of opposition shall be filed in a written reasoned statement. It shall not be deemed to have been filed until the opposition fee has been paid (Art. 99(1) European patent convention).

Description

This invention relates to a method of encapsulation, in particular of encapsulating hydrophobic liquids, and to encapsulated products containing such liquids. It is concerned particularly, although not necessarily exclusively, with encapsulated products that are to be used as additions, eg. for fragrance or flavour, to a base product.

There are many known micro-encapsulation methods which employ silica particles as an encapsulating material, the function of which is to allow a delayed but continuous release of the material encapsulated. In one type, exemplified by US 4440542 and US 4579779, the silica particles form an open-pored matrix which holds the material, especially liquids, in its interstices. In such a structure the outer pores are exposed and the encapsulated material may not be adequately shielded from premature release, whether by chemical reaction or by leaching. Furthermore, these porous structures rely on physical sorption of the encapsulated material and have a relatively limited liquid-silica ratio, or loading. That disadvantage is more pronounced if the encapsulated material is prone to premature release since this would be countered by increasing the density of the porous structure.

These porous structures can be produced in a modified form with a substantially impervious outer shell, as exemplified in US 4464317. Although this prevents leaching of the encapsulated material it reduces still further the loading that can be achieved. A further disadvantage of the processes described in US 4464317 is that they produce relatively large particles (eg. $>500\mu\text{m}$), of a size that is unsuitable for many processes.

In another type of process described in EP294206A, porous spheroidal silica particles are formed which contain dispersed droplets of an encapsulated liquid. It is said that a loading can be achieved of up to 50% by weight of the silica, which is not a remarkable figure, yet the dry product still has considerable porosity since, when encapsulating a perfume, it is said that any wetting produces an appreciable increase of odour. Furthermore, the preparation of the product is complex and lengthy.

In another process described in EP309054A an aqueous emulsion is formed in which droplets of lubricating oil are coated with silica particles which stabilise the emulsion. The oil is added to a partially hydrophobed silica dispersion prepared from colloidal silica and the mixture is homogenised in a high-shear mixer. In the resulting emulsion the oil droplets are held as silica-coated droplets that are stable until ruptured by pressure, as when a lubricating surface, to produce a lubricating film.

Experiments have shown, however, that if such emulsions are formed using hydrophobic materials, such as fragrances and flavours, that do not dissolve unaided in water, these emulsions cannot be usefully employed as additives to consumer products. This is because the emulsified material can make contact with and be dissolved by other components such as surfactants in cleaning or washing products or mineral oils in cosmetic products. The emulsified material will thus simply disperse when the silica-coated droplets are mixed into the product.

According to one aspect of the present invention, there is provided a process for encapsulating a hydrophobic material in which an aqueous dispersion of silica of particle size not substantially greater than 100nm is produced, an emulsion is formed by high-shear mixing of said dispersion with the material to be encapsulated to give a weight ratio of encapsulated material to silica of at least 1.5:1, said emulsion being stabilized by the addition of a gelling agent.

Observations indicate that the gelling step forms or stabilises a shell-like structure of silica particles around individual droplets of the hydrophobic material in the emulsion. The resulting capsules have sufficient stability to be used as such in certain applications. It is found, furthermore, that it is possible to give such shell-like structures significant strength and a good degree of imperviousness in relation to the encapsulated material.

The shell-like encapsulating structures formed are also found to be capable of holding relatively high loadings of hydrophobic material and the invention can give a loading of the encapsulated hydrophobic material to the outer silica layer greater than 2:1, eg. up to 100:1.

Typical products in which the micro-encapsulates can be incorporated include cosmetic products, such as skin cream and sunscreen formulations, detergent products such as laundry wash products, household cleaners, shampoos, hair conditioners and bleaches, and oral hygiene products such as toothpastes. Depending upon the product and its use, the encapsulation may be employed to protect materials against loss by evaporation during storage or hot processing, or against chemical degradation (particularly in products containing other, hostile ingredients), to improve the targeting of materials in use (eg. perfume deposition onto fabrics during washing and conditioning) particularly to assist triggered delivery through heat, shear or dissolution, or to extend activity (eg. of a fragrance or flavouring) through controlled delivery and evaporation.

Suitable and convenient silica materials include colloidal silica and fumed silica. Colloidal silica is supplied commercially as an alkaline solution (pH

>9) and a hydrophobing agent may be added to help produce the aqueous dispersion. Suitable hydrophobing agents include positively charged surfactants such as cetyltrimethyl-ammonium bromide (CTAB). Fumed silica forms an acidic emulsion (pH 3-5) which requires no hydrophobing agent.

By using silica particles having some hydrophobic character in the aqueous dispersion, or by giving them such a character by adding a hydrophobing agent, the particles tend to concentrate at the interface between the water and the hydrophobic material as high-shear mixing occurs and so coat the (dispersed) hydrophobic material. It is found possible, in fact, to obtain silica-stabilised emulsions having relatively high ratios of emulsified material to silica, with the weight of emulsified hydrophobic material at least equal to the weight of silica. At this stage, however, the silica particles are bound together too weakly to hold a hydrophobic material stably when incorporated in a product.

The porosity of the initial coating of silica particles is related to some extent to the silica particle size which may be up to 100nm. By using silica with a relatively small particle size, preferably in the range 5-50nm, or more advantageously 5-20nm, a shell-like structure is obtained which can be strengthened and rendered substantially impervious after emulsification. Mixtures of silica particles of different sizes can also be used to improve packing and so improve further the stability of the shell-like structure.

The treatment after emulsification comprises a gelling step. It may be sufficient to perform that step using an appropriate gelling agent, but otherwise the gelling step can be followed with a further treatment.

In the first case, the gelling step can be carried out in a coacervation process using a positively charged compound such as a positively charged polymer or a gum or a silicone which, as it attaches itself to the silica shell, is able to form an outer layer on the capsule.

Alternatively gelling can be obtained by acidifying the emulsion to a pH value of approximately 8.0 to 5.0, preferably 6.5 to 5.5 by the addition of an acid or by changing the charge state of the silica shell by adding a surfactant or a salt such as CaCl_2 . With this procedure, the product may be useful for limited applications, eg. as a slow-release air freshener or carpet freshener or deodorant. However, for most purposes further treatment may be needed to reduce the porosity of the silica shell and/or to otherwise modify the effective retention of the encapsulated product, eg. to increase the strength of the capsules against rupture.

There are a number of alternative treatments available for reducing the porosity of the capsules in the gelled product. The choice may depend on the form (eg. liquid or solid) of final product required. For example, by heating or adding a drying agent after gelling, a dried product can be obtained that is suitable for use in powder formulations, eg. to carry a fragrance in an air freshener or a laundry powder.

In another form of treatment, the capsules can be given an outer coating by blending the gelled material with a suitable material such as a liquid crystal forming surfactant or a polymer. Suitable surfactants may be selected from any of the major classes (ie. non-ionic, cationic, anionic and zwitterionic) and where appropriate their ability to form liquid crystal structures may be assisted by the inclusion of structuring aids such as steroids. Examples of suitable surfactants include

Polyethoxylated fatty alcohols (eg. POE(2) cetyl alcohol)
Sorbitan esters (eg. sorbitan monostearate)
Glycerol esters (eg. glycerol monostearate)
Phospholipids (eg. phosphatidylcholine)
Fatty alcohols (eg. cetyl alcohol)
Quaternaries (eg. dimethylditallowammonium chloride)

Suitable polymers may include starch, modified starch, other polysaccharides (eg. gums), derivatised polysaccharides and synthetic polymers such as silicones, polyacrylics (eg. polyacrylamide) and polyvinyl pyrrolidones.

When treated with a liquid crystal forming surfactant, in particular, the capsules may be given a greater resistance to rupture, especially against abrasion, which can improve their efficacy for products such as toothpaste flavourings or sunscreen formulations. Addition of starch may be appropriate for a fragrance, eg. in a carpet freshener formulation when some of the fragrance will be held also by the starch as it precipitates out of solution while the product dries and will be released slowly after the carpet has dried. Preferably, with capsules such as these, having a further coating material, the encapsulated material forms at least 60% by weight of the encapsulate and may be present in up to 30 times the weight of the encapsulating material.

Micro-encapsulates prepared according to the invention can be incorporated in final products in any required proportion up to 99%, but for most purposes a minimum level of 0.1-0.2% is appropriate, in particular if the product also has a further amount of the hydrophobic material freely dispersed in it.

The invention is illustrated by the following examples of some processes and products according to the invention. The first group of Examples 1-

4 d scribe the formation of a preliminary emulsion with the silica particles.

EXAMPLE 1

0.01g CTAB was dissolved in 75g de-ionised water. 25g colloidal silica (Ludox HS-40 from E I Du Pont de Nemours, mean particle size 12nm) was then mixed with the water to form a homogeneous dispersion for encapsulating a perfume having the formulation: coumarin 0.5%, benzyl acetate extra 4.0%, benzyl salicylate 10.0%, dihydromyrcenol 10.0%, citronellol 10.0%, Lixetone (TM) 8.0%, methyl dihydrojasmonate 5.0%, phenyl ethyl alcohol 10.0%, Traseolide (TM) 7.5%, Jas-mopyrane (TM) forte 10.0%, linalol 10.0%, hexyl cinnamic aldehyde 8.0%, isolongifolanone 3.0%, styrallyl acetate 1.0%, methylionone 3.0%. 75g of the perfume was added to the dispersion while mixing in a high shear mixer, eg. a Silverson mixer, for approximately 30 seconds. A milky-white emulsion resulted with a pH of approximately 9 in which droplets of the perfume were encapsulated by the silica. The capsule size was found to be in the range 2-5µm and the weight ratio of silica to encapsulated material was 1:7.5.

EXAMPLE 2

Example 1 was repeated using 78g de-ionised water, without CTAB, 2g fumed silica (Aerosil 300 from Degussa AG, mean particle size 7nm) with 20g flavouring as the hydrophobic material. The formulation of the flavouring was: anethole 7.06% carvone laevo 8.90%, menthol laevo 10.0%, peppermint american 23.0%, spearmint american 51.0%, vanillin 0.04%. With the high-shear mixing a milky-white emulsion was produced with a pH of approximately 3.5. The particle size was in the range 2-30µm and the weight ratio of silica to encapsulated material was 1:10.

EXAMPLE 3

An aqueous dispersion was prepared using 15g Ludox HS-40 and 5g Ludox SM, particle size 7nm in 60g de-ionised water. The dispersion was mixed in a high shear mixer with 20g parsol MCX (ex Givaudan) a sunscreen active, to give a milky-white emulsion with a pH of approximately 10. The particle size was in the ratio 5-10µm and the weight ratio of silica to encapsulated material was 1:2.66.

EXAMPLE 4

Using the procedure of Example 1 a homogeneous dispersion was produced from 59.5g de-ionised water and 0.5g Aerosil 200, particle size 12nm.

This was mixed with 30g of a perfume having the formulation: Galaxolide (TM) 7.5%, Lixetone (TM) 12.0%, linalol 7.0%, benzyl salicylate 7.2%, hexyl cinnamic aldehyde 13.0%, methyl dihydrojasmonate 14.6%, phenyl ethyl alcohol 9.0%, dipropylene glycol 18.0%, litsea cubeba 5.0%, coumarin 0.06%, linalol 3.6%, hexyl salicylate 3.04%. A milky-white emulsion resulted with a pH of approximately 3.5. The particle size was 2-10µm and the weight ratio of silica to encapsulated material was 1:60.

The emulsions obtained in these examples showed themselves to be unstable when added to a product base, the silica coatings being quite porous. As a consequence the encapsulated material eventually dispersed, eg. dissolving in an aqueous base. In the following group of Examples 5-9 the emulsion was acidified in a gelling operation giving a micro-porous silica shell structure. Porosity was reduced, giving products suitable for some end uses, as already mentioned, but generally further processing was desirable for completion of the encapsulation.

EXAMPLE 5

The pH of the emulsion of Example 1 was adjusted to 6.5 using 0.5M HCl (aqueous) and was allowed to stand for 72 hours. A thick cream was formed which "broke" on shaking to give a fluid emulsion.

EXAMPLE 6

The pH of the emulsion of Example 3 was adjusted to 6.0 using 0.5M HCl (aqueous) and the emulsion was allowed to stand for 48 hours. This produced a thick cream which "broke" on shaking to give a fluid emulsion. The pH was then increased to 7.5 to assist stability of the emulsion.

EXAMPLE 7

5g of 0.1M CaCl₂ (aqueous) was added to 100g of the emulsion of Example 1. Gelation took place over the course of several days.

EXAMPLE 8

10g of 0.1M CaCl₂ (aqueous) was added to 100g of the emulsion of Example 2. Rapid gelling followed to give a thick creamy product.

EXAMPLE 9

The pH of Example 4 was adjusted to 6.5 with 0.5M HCl (aqueous) and then 10g of 5% w/w CaCl₂ in aqueous solution was added to 50g of the emulsion. Rapid gelation followed.

The following group of Examples 10-15 illustrate final treatments for gelled intermediate products such as are produced by the preceding group of Examples 5-9 to reduce their porosity or otherwise to adapt them better to a particle end use.

EXAMPLE 10

A mixture of 4g stearyl alcohol 1.5g sorbitan monostearate - Span 60 (ICI Speciality Chemicals) and 0.25g sodium lauryl sulphate were blended together by heating to form a homogeneous liquid. This mixture of the three surfactants was then added to 50g of the gelled product of Example 7 which had been heated to the same temperature, while stirring gently. Stirring was continued while cooling. The product was a creamy stable emulsion suitable for incorporation into an aqueous detergent product.

EXAMPLE 11

5g starch in the form of Capsul (National Starch) was dissolved in 40g de-ionised water. The solution was dispersed in 50g of the gelled product of Example 5. The resulting product was suitable for a carpet freshening product.

EXAMPLE 12

50g of the gelled product of Example 5 was dehydrated with 50g sodium tripolyphosphate to give a dry powder holding a fragrance that could be used in a laundry powder.

EXAMPLE 13

The gelled product of Example 5 was air-dried to give a dry powder containing 60% w/w perfume that could be used in a laundry powder.

EXAMPLE 14

50g of the gelled product of Example 9 was prepared but with the flavouring described in Example 2 as the encapsulated material. This gelled product was further processed by the method of Example 10 using 4.5g glycerol monostearate, 0.5g CTAB. The final product was suitable for flavouring toothpaste.

EXAMPLE 15

The method of Example 14 was followed using 50g of the gelled product of Example 6, 4.5g arachidyl alcohol and 0.5g CTAB, to produce an encapsulate carrying a sunscreen agent.

EXAMPLE 16

Following the method of Example 10, 50g of the gelled product of Example 9 was mixed with 4.5g POE(2) cetyl alcohol Brij 52 (ICI Speciality Chemicals) and 0.5g sodium lauryl sulphate. The product was a stable emulsion suitable for perfuming a hard-surface cleaner.

The initial emulsions such as those in Examples 1-4 can alternatively be gelled in a complex coacervation process to reduce the porosity of the silica, even to the extent of rendering the shell substantially impervious, and increase the strength of the encapsulation. This is illustrated in the following Examples 17-21.

EXAMPLE 17

The pH of the emulsion of Example 2 was adjusted to 7.5 using 1M HCl(aqueous). After standing for 6 hours, 1g of gum acacia in a 10% aqueous solution was then added to 100g of the emulsion. The product was suitable for inclusion in a dental care formulation.

EXAMPLE 18

The pH of the emulsion of Example 3 was adjusted to 7.0 using 0.5M HCl(aqueous). 2g of quaternised starch derivative Celquat L-200 (National Starch Inc) in a 5% aqueous solution was added to 60g of the emulsion. A thick creamy product resulted suitable for incorporation in a sunscreen preparation.

EXAMPLE 19

The pH of an emulsion prepared according to the method of Example 1 but using 37.5g perfume was adjusted to 7.5 using 0.5M HCl(aqueous). 2g of cationic silicone Abilquat 3270 (Th. Goldschmidt AG) was added to 75g of the emulsion. The product was a stable emulsion suitable for inclusion in a hair conditioner.

EXAMPLE 20

To the product of Example 1 was added 5% w/w of Crystal 100 (Crosfield Chemicals Ltd), a sodium silicate formulation, while simultaneously adjusting the pH to and maintaining it at about 7. 2% w/w of CaCl_2 in a 0.5M aqueous solution was added to gel the sodium silicate around the emulsion droplets. An improved encapsulate resulted.

EXAMPLE 21

To the product of Example 2 was added 5% w/w of Crystal 100 (Crosfield Chemicals Ltd), a sodium

silicate formulation, while simultaneously adjusting the pH to and maintaining it at about 7.2% w/w of Celquat (National Starch Inc) in a 5% aqueous solution was added to gel the sodium silicate around the emulsion droplets. The product was suitable for flavouring an oral hygiene product.

Examples of the incorporation of the encapsulated materials of the preceding examples in aqueous base products will now be given.

EXAMPLE 22

A toothpaste was produced by first formulating three phases as follows, where the proportions are given as percentages by weight of the finished product:

- (A) sorbitol (70%aq) 30.0%, sodium monofluorophosphate 0.85%, water 38.95%
- (B) syloblanc 81 (silica) (W.R Grace & Co) 20.0%, sodium carboxymethylcellulose 1.60%, titanium dioxide 0.50%
- (C) sorbitol (70%aq) 5.0%, NaOH (50%aq) 0.40%, empicol LXV (Albright and Wilson) 1.50%, saccharin 0.20%.

Phases A and B were mixed thoroughly under high-shear for 30 minutes. Phase C was then stirred in under vacuum while maintaining the temperature at 40°C. Finally 1% flavour in the form of an encapsulated product containing a flavouring agent was mixed in under low shear for 15 minutes.

To test the comparative efficacy of the encapsulate stabilising measures employed in accordance with the invention three toothpaste samples were prepared each with the products of one of Examples 2, 8 and 14 added to the base, and a fourth sample was prepared with flavouring agent treated in accordance with Example 14 but without previously producing an emulsion of the agent with silica particles. A panel of trained evaluators assessed the products and reported a substantially greater flavour impact with the sample containing the silica-encapsulated product of Example 14 indicating that in comparison with all of the other samples the capsules had retained substantially more of the flavouring to release it upon use.

EXAMPLE 23

A skin cream was produced, first formulating two phases

- (A) Emulgin B1 (Henkel Chemicals Ltd) 3.0%, Eutanol G (Henkel Chemicals Ltd) 22.0%, isopropyl myristate 5.0%, glycerol monostearate 15.0%
- (B) Hygroplex HHG (Henkel Chemicals Ltd) 3.0%, preservative q.s., purified water 51.8%.

Each phase was heated separately to 70°C. Phase B was then mixed into phase A with constant

stirring which continued until the resulting cream was cool. To this was then added 1% of an encapsulation product containing a perfume.

The skin cream base was used for four test samples containing perfume encapsulated by the processes of Examples 1, 7, 10 and 20 respectively. Examination by microscope showed that the samples using the encapsulates of Examples 1 and 7 were unstable, the capsules simply dispersing in the first and the perfume leaching out in the second, whereas the products of Examples 10 and 20 were stable in the product.

It will be understood that these and other products incorporating micro-encapsulates prepared in accordance with the invention may include free further hydrophobic compounds as fragrances, flavourings etc.

Claims

1. A process for encapsulating a hydrophobic material in which an aqueous dispersion of silica of a particle size not substantially greater than 100nm is produced, an emulsion then being formed by high-shear mixing of said dispersion with the material to be encapsulated, the weight ratio of encapsulated material to silica being at least 1.5:1, and said emulsion being stabilized by the addition of a gelling agent effective to form or stabilise an encapsulating structure of silica particles around droplets of the hydrophobic material.
2. A process according to claim 1 wherein the hydrophobic material is a flavour, a fragrance or a sunscreen active.
3. A process according to claim 1 or claim 2 wherein the silica particles are in the form of colloidal silica or fumed silica having a size not substantially greater than 50nm, and is preferably between 5 and 20nm.
4. A process according to claim 1, claim 2 or claim 3 wherein the weight ratio of encapsulated material to silica is greater than 2:1.
5. A process according to any one of the preceding claims wherein the gelling step comprises acidifying the emulsion and/or adding a salt and/or a positively charged surfactant.
6. A process according to any one of the preceding claims wherein the gelled product is dried.
7. A process according to any one of claims 1 to 6 wherein a further material is coated onto the capsules to reduce their porosity.

8. A process according to claim 5 together with claim 7 wherein said further material is a liquid-crystal-forming surfactant or starch or other polymer.

9. A process according to any one of claims 1 to 3 wherein the emulsion is gelled by the addition of a positively charged material that forms an outer coating on the capsules.

10. A process according to claim 9 wherein said material is a positively charged polymer and/or a gum and/or a silicone.

11. A process according to any one of claims 7 to 10 wherein the encapsulated material forms at least 60% of the weight of the capsules.

12. A micro-encapsulate comprising a gelled outer silica layer containing a hydrophobic material in which the loading of said material to the silica is at least 1.5:1.

13. A micro-encapsulate according to claim 12 in the form of a dried particulate.

14. A product comprising the micro-encapsulate of claim 12 as an additive in a liquid base.

15. A product according to claim 13 or claim 14 wherein the micro-encapsulate provides a flavour and/or fragrance and/or cosmetic ingredient.

16. A cosmetic product comprising at least 0.02% by weight of a micro-encapsulate prepared according to any one of claims 1 to 10 wherein the hydrophobic material is a perfume and/or a cosmetic ingredient.

17. A product according to claim 16 formulated as a skin cream.

18. A product according to claim 15, claim 16 or claim 17 wherein the cosmetic ingredient is a sunscreen active.

19. A detergent product comprising at least 0.02% by weight of a micro-encapsulate prepared according to any one of claims 1 to 10 wherein the hydrophobic material is a perfume.

20. A product according to claim 19 formulated as a laundry wash product or a household cleaner or a bleach or a shampoo or hair conditioner.

21. An oral hygiene product comprising at least 0.01% by weight of a micro-encapsulate prepared

according to any one of claims 1 to 11 wherein the hydrophobic material is a flavour and/or a food ingredient and/or a therapeutic agent.

5

Patentansprüche

10

1. Verfahren zur Verkapselung eines hydrophoben Materials, bei dem eine wäßrige Dispersion von Siliziumdioxid mit einer Teilchengröße von nicht wesentlich größer als 100nm hergestellt wird, anschließend durch stark scherenendes Vermischen der Dispersion mit dem zu verkapselnden Material eine Emulsion gebildet wird, wobei das Gewichtsverhältnis des verkapselten Materials zum Siliziumdioxid mindestens 1,5 : 1 beträgt, und die Emulsion durch Zugabe eines Geliermittels stabilisiert wird, welches zur Bildung oder Stabilisierung einer verkapselnden Struktur aus Siliziumdioxid-Teilchen um die Tröpfchen des hydrophoben Materials geeignet ist.

15

20

25

2. Verfahren nach Anspruch 1, worin das hydrophobe Material ein Aromastoff, ein Duftstoff oder ein Lichtschutzwirkstoff ist.

30

3. Verfahren nach Anspruch 1 oder 2, worin die Siliziumdioxid-Teilchen in Form von kolloidalem Siliziumdioxid oder Kieselstaub mit einer Größe vorliegen, die nicht wesentlich größer als 50nm ist und vorzugsweise zwischen 5 und 20nm liegt.

35

40

4. Verfahren nach Anspruch 1, 2 oder 3, worin das Gewichtsverhältnis des verkapselten Materials zum Siliziumdioxid größer als 2:1 ist.

45

5. Verfahren nach einem der vorstehenden Ansprüche, worin der Gelierschritt das Ansäuern der Emulsion und/oder den Zusatz eines Salzes und/oder eines positiv geladenen oberflächenaktiven Mittels umfaßt.

50

6. Verfahren nach einem der vorstehenden Ansprüche, worin das gelierte Produkt getrocknet wird.

55

7. Verfahren nach einem der Ansprüche 1 bis 6, worin ein weiteres Material auf die Kapseln aufgebracht wird, um deren Porosität zu reduzieren.

8. Verfahren nach Anspruch 5 und 7 zusammengefaßt, worin das weitere Material ein Flüssigkristall-bildendes oberflächenaktives Mittel, eine Stärke oder ein weiteres Polymer ist.

9. Verfahren nach einem der Ansprüche 1 bis 3, worin die Emulsion durch die Zugabe eines positiv geladenen Materials geliert wird, welches eine äußere Hülle auf den Kapseln bildet.

10. Verfahren nach Anspruch 9, worin das Material ein positiv geladenes Polymer und/oder ein Gum und/oder ein Silicon ist.

11. Verfahren nach einem der Ansprüche 7 bis 10, worin das verkapselte Material mindestens 60% des Gewichts der Kapseln darstellt.

12. Mikrokapseln mit einer äußeren Schicht aus geliertem Siliziumdioxid, enthaltend ein hydrophobes Material, in denen die Füllung des besagten Materials zum Siliziumdioxid mindestens 1,5:1 beträgt.

13. Mikrokapseln nach Anspruch 12 in Form getrockneter Feststoffteilchen.

14. Produkt, welches die Mikrokapseln nach Anspruch 12 als ein Additiv in einer flüssigen Grundlage umfaßt.

15. Produkt nach Anspruch 13 oder 14, worin die Mikrokapseln als ein Aromastoff und/oder ein Duftstoff und/oder ein kosmetischer Inhaltsstoff dienen.

16. Kosmetisches Mittel, welches mindestens 0,02 Gew.-% nach Ansprüche 1 bis 10 hergestellte Mikrokapseln umfaßt, worin das hydrophobe Material ein Parfüm und/oder ein kosmetischer Inhaltsstoff ist.

17. Produkt nach Anspruch 16, formuliert als eine Hautcreme.

18. Produkt nach Anspruch 15, 16 oder 17, worin der kosmetische Inhaltsstoff ein Lichtschutzwirkstoff ist.

19. Reinigungsmittel, welches mindestens 0,02 Gew.-% nach einem der Ansprüche 1 bis 10 hergestellte Mikrokapseln umfaßt, worin das hydrophobe Material ein Parfüm ist.

20. Produkt nach Anspruch 19, formuliert als ein Waschmittel, ein Haushaltsreiniger, ein Bleichmittel, ein Shampoo oder ein Haarkonditionierungsmittel.

21. Mundhygieneprodukt, welches mindestens 0,01 Gew.-% nach einem der Ansprüche 1 bis 11 hergestellte Mikrokapseln umfaßt, worin das hydrophobe Material ein Aromastoff und/oder

ein Nahrungsmittelzusatzstoff und/oder ein Heilmittel ist.

Revendications

1. Procédé d'encapsulation d'une matière hydrophobe dans lequel une dispersion aqueuse de silice d'une granulométrie ne dépassant pas sensiblement 100 nm est produite, une émulsion étant ensuite formée par mélange par cisaillement intime de ladite dispersion avec la matière à encapsuler, le rapport pondéral de la matière encapsulée sur la silice étant d'au moins 1,5/1, et ladite émulsion étant finalement stabilisée par l'addition d'un agent gélifiant efficace pour former ou stabiliser une structure encapsulante de particules de silice enrobant des gouttelettes de la matière hydrophobe.

2. Procédé selon la revendication 1 dans lequel la matière hydrophobe est un arôme, un parfum ou un principe actif formant écran solaire.

3. Procédé selon la revendication 1 ou la revendication 2 dans lequel les particules de silice se présentent sous la forme de silice colloïdale ou de silice fumée d'une taille qui n'est pas sensiblement supérieure à 50 nm, et de préférence comprise entre 5 et 20 nm.

4. Procédé selon la revendication 1, la revendication 2 ou la revendication 3 dans lequel le rapport pondéral de la matière encapsulée sur la silice est supérieur à 2/1.

5. Procédé selon l'une des revendications précédentes dans lequel l'étape de gélification comprend l'acidification de l'émulsion et/ou l'addition d'un sel et/ou d'un agent tensioactif chargé positivement.

6. Procédé selon l'une des revendications précédentes dans lequel le produit gélifié est séché.

7. Procédé selon l'une des revendications 1 à 6 dans lequel les capsules sont enduites d'une matière supplémentaire pour réduire leur porosité.

8. Procédé selon la revendication 5 et la revendication 7 dans lequel ladite matière supplémentaire est un agent tensioactif, un amidon ou tout autre polymère formant un liquide cristallin.

9. Procédé selon l'une des revendications 1 à 3 dans lequel l'émulsion est gélifiée par addition d'une matière chargée positivement qui form

un revêtement externe sur les capsules.

10. Procédé selon la revendication 9 dans lequel ladite matière est un polymère et/ou une gomme et/ou un silicone chargés positivement. 5
11. Procédé selon l'une des revendications 7 à 10 dans lequel la matière encapsulée représente au moins 60% en poids des capsules. 10
12. Microcapsules comprenant une couche externe de silice gélifiée contenant une matière hydrophobe dans lesquelles la charge de ladite matière hydrophobe dans la silice est d'au moins 1,5/1. 15
13. Microcapsules selon la revendication 12 sous la forme d'une particule séchée.
14. Produit comprenant les microcapsules de la revendication 12 en tant qu'additif dans une base liquide. 20
15. Produit selon la revendication 13 ou la revendication 14 dans lequel les microcapsules contiennent un arôme et/ou un parfum et/ou un ingrédient cosmétique. 25
16. Produit cosmétique comprenant au moins 0,02% en poids des microcapsules préparées selon l'une des revendications 1 à 10 dans lequel la matière hydrophobe est un parfum et/ou un ingrédient cosmétique. 30
17. Produit selon la revendication 16 formulé en tant que crème cutanée. 35
18. Produit selon la revendication 15, la revendication 16 ou la revendication 17 dans lequel l'ingrédient cosmétique est un principe actif formant écran solaire. 40
19. Produit détergent comprenant au moins 0,02% en poids des microcapsules préparées selon l'une des revendications 1 à 10 dans lequel la matière hydrophobe est un parfum. 45
20. Produit selon la revendication 19 formulé en tant que produit de lessive ou nettoyant ménager ou décolorant ou shampoing ou lotion capillaire. 50
21. Produit d'hygiène orale comprenant au moins 0,01% en poids des microcapsules préparées selon l'une des revendications 1 à 11 dans lequel la matière hydrophobe est un arôme et/ou un ingrédient alimentaire et/ou un agent thérapeutique. 55